



## SGSH gene

N-sulfoglucosamine sulfohydrolase

### Normal Function

The *SGSH* gene provides instructions for producing an enzyme called sulfamidase. This enzyme is located in lysosomes, compartments within cells that digest and recycle different types of molecules. Sulfamidase is involved in the step-wise breakdown of large molecules called glycosaminoglycans (GAGs). GAGs are composed of sugar molecules that are linked together to form a long string. To break down these large molecules, individual sugars are removed one at a time from one end of the molecule. Sulfamidase removes a chemical group known as a sulfate from a sugar called glucosamine when it is at the end of the GAG chain.

### Health Conditions Related to Genetic Changes

#### mucopolysaccharidosis type III

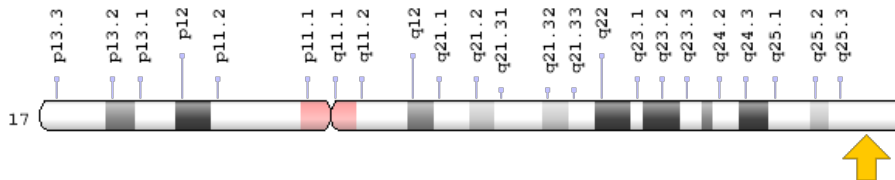
More than 80 mutations in the *SGSH* gene have been found to cause mucopolysaccharidosis type IIIA (MPS IIIA). Most of these mutations change single DNA building blocks (nucleotides) in the gene. All of the mutations that cause MPS IIIA reduce or eliminate the function of sulfamidase.

The lack of sulfamidase activity disrupts the breakdown of a subset of GAGs called heparan sulfate. As a result, partially broken down heparan sulfate accumulates within lysosomes. Researchers believe that the accumulation of GAGs interferes with the functions of other proteins inside the lysosomes and disrupts the normal functions of cells. It is unknown why the buildup of heparan sulfate mostly affects the central nervous system in MPS IIIA.

## Chromosomal Location

Cytogenetic Location: 17q25.3, which is the long (q) arm of chromosome 17 at position 25.3

Molecular Location: base pairs 80,200,668 to 80,220,400 on chromosome 17 (Homo sapiens Annotation Release 108, GRCh38.p7) (NCBI)



Credit: Genome Decoration Page/NCBI

## Other Names for This Gene

- heparan N-sulfatase
- heparan sulfate sulfatase
- HSS
- N-sulphoglucosamine sulphohydrolase
- N-sulphoglucosamine sulphohydrolase precursor
- SFMD
- SPHM\_HUMAN
- sulfamidase
- sulfoglucosamine sulfamidase
- sulphamidase

## Additional Information & Resources

### Educational Resources

- Eureka Bioscience Collection: Defects in Glycosaminoglycan Degradation (Mucopolysaccharidoses)  
<https://www.ncbi.nlm.nih.gov/books/NBK6177/#A53462>

### Scientific Articles on PubMed

- PubMed  
<https://www.ncbi.nlm.nih.gov/pubmed?term=%28%28SGSH%5BTIAB%5D%29+OR+%28N-sulfo-glucosamine+sulfohydrolase%5BTIAB%5D%29%29+OR+%28%28heparan+sulfate+sulfatase%5BTIAB%5D%29+OR+%28MPS3A%5BTIAB%5D%29+OR+%28N-sulphoglucosamine+sulphohydrolase%5BTIAB%5D%29+OR+%28sulphamidase%5BTIAB%5D%29+OR+%28heparan+N-sulfatase%5BTIAB%5D%29+OR+%28sulfamidase%5BTIAB%5D%29%29+AND+%28%28Genes%5BMH%5D%29+OR+%28Genetic+Phenomena%5BMH%5D%29%29+AND+human%5Bmh%5D+AND+%22last+3600+days%22%5Bdp%5D>

### OMIM

- N-SULFOGLUCOSAMINE SULFOHYDROLASE  
<http://omim.org/entry/605270>

### Research Resources

- ClinVar  
<https://www.ncbi.nlm.nih.gov/clinvar?term=SGSH%5Bgene%5D>
- HGNC Gene Family: Sulfatases  
<http://www.genenames.org/cgi-bin/genefamilies/set/410>
- HGNC Gene Symbol Report  
[http://www.genenames.org/cgi-bin/gene\\_symbol\\_report?q=data/hgnc\\_data.php&hgnc\\_id=10818](http://www.genenames.org/cgi-bin/gene_symbol_report?q=data/hgnc_data.php&hgnc_id=10818)
- NCBI Gene  
<https://www.ncbi.nlm.nih.gov/gene/6448>
- UniProt  
<http://www.uniprot.org/uniprot/P51688>

### **Sources for This Summary**

- Meyer A, Kossow K, Gal A, Steglich C, Mühlhausen C, Ullrich K, Bräulke T, Muschol N. The mutation p.Ser298Pro in the sulphamidase gene (SGSH) is associated with a slowly progressive clinical phenotype in mucopolysaccharidosis type IIIA (Sanfilippo A syndrome). *Hum Mutat.* 2008 May;29(5):770. doi: 10.1002/humu.20738.  
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